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## UNITED STATES EPARTMENT OF COMMERCE United States Pat int and Trad mark Offic

Address: COMMISSIONER OF PATENTS AND TRADEMARKS Washington, D.C. 20231

APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR			ATTORNEY DOCKET NO.	
09/202,463	08/19/99	BRUNDELL		J	100096.401	
Г		HM22/0820	7	EXAMINER		
SEED & BERRY 6300 COLUMBIA CENTER		HI1227 0020		TURŅER.	s	
				ART UNIT ,	PAPER NUMBER	
SEATTLE WA	98104-7092			1647	<u>(</u> 0	
		•	;	DATE MAILED:	08/20/01	

Please find below and/or attached an Office communication concerning this application or proceeding.

**Commissioner of Patents and Trademarks** 

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### Office Action Summary

Application No. 09/202,463

Appl ....(s)

Brundell et al.

Examiner

Sharon L. Turner, Ph.D.

Art Unit 1647



The MAILING DATE of this communication appear	s on the cover sheet with the correspondence address
Period for Reply  A SHORTENED STATUTORY PERIOD FOR REPLY IS SETHE MAILING DATE OF THIS COMMUNICATION.  - Extensions of time may be available under the provisions of 37 CFR 1. after SIX (6) MONTHS from the mailing date of this communication.  - If the period for reply specified above is less than thirty (30) days, a replace to the considered timely.  - If NO period for reply is specified above, the maximum statutory period communication.  - Failure to reply within the set or extended period for reply will, by statut.  - Any reply received by the Office later than three months after the mailing earned patent term adjustment. See 37 CFR 1.704(b).  Status	136 (a). In no event, however, may a reply be timely filed  bly within the statutory minimum of thirty (30) days will  will apply and will expire SIX (6) MONTHS from the mailing date of this  e, cause the application to become ABANDONED (35 U.S.C. § 133).
1) X Responsive to communication(s) filed on	
2a) ☐ This action is <b>FINAL</b> . 2b) ☒ This act	ion is non-final.
3) Since this application is in condition for allowance e closed in accordance with the practice under Exp	
Disposition of Claims	
4) ☒ Claim(s) <u>1-18</u>	is/are pending in the applica
4a) Of the above, claim(s)	is/are withdrawn from considera
5) Claim(s)	is/are allowed.
6)  Claim(s)	is/are rejected.
•	is/are objected to.
8) 🗓 Claims <u>1-18</u>	are subject to restriction and/or election requirem
Application Papers  9) ☐ The specification is objected to by the Examiner.  10) ☐ The drawing(s) filed on is/a  11) ☐ The proposed drawing correction filed on	
12) The oath or declaration is objected to by the Examin	er.
Copies of the certified copies of the priority documents application from the International Bureau *See the attached detailed Office action for a list of the	been received.  been received in Application No  cuments have been received in this National Stage (PCT Rule 17.2(a)).  certified copies not received.
14) Acknowledgement is made of a claim for domestic p	ononly under 35 U.S.C. 9 T19(e).
Attachment(s)	
15) Notice of References Cited (PTO-892)	18) Interview Summary (PTO-413) Paper No(s).
<ul> <li>16) Notice of Draftsperson's Patent Drawing Review (PTO-948)</li> <li>17) Information Disclosure Statement(s) (PTO-1449) Paper No(s).</li> </ul>	19) Notice of Informal Patent Application (PTO-152)  20) Other:

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#### **DETAILED ACTION**

#### Sequence Compliance

1. This application contains sequence disclosures that are encompassed by the definitions for nucleotide and/or amino acid sequences set forth in 37 CFR 1.821(a)(1) and (a)(2). However, this application fails to comply with the requirements of 37 CFR 1.821 through 1.825 for the reason(s) set forth on the attached Notice To Comply With Requirements For Patent Applications Containing Nucleotide Sequence And/Or Amino Acid Sequence Disclosures. Applicant must comply with the requirements of the sequence rules (37 CFR 1.821 - 1.825) before the application can be examined under 35 U.S.C. §§ 131 and 132.

Applicant is given ONE MONTH from the mailing date of this communication within which to comply with the sequence rules, 37 CFR 1.821 - 1.825. Failure to comply with these requirements will result in ABANDONMENT of the application under 37 CFR 1.821(g). Extensions of time may be obtained by filing a petition accompanied by the extension fee under the provisions of 37 CFR 1.136(a). Direct the reply to the undersigned. Applicant is requested to return a copy of the attached Notice to Comply with the reply.

In particular applicant is directed to MPEP 2423, 37 CFR 1.822(e) and claim 6 which does not comply with the sequence rules.

#### Election/Restrictions

2. Restriction is required under 35 U.S.C. 121 and 372.

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This application contains the following inventions or groups of inventions which are not so linked as to form a single general inventive concept under PCT Rule 13.1.

In accordance with 37 CFR 1.499, applicant is required, in response to this action, to elect a single invention to which the claims must be restricted.

Groups I-V, claim(s) 1-6, 11 and 16-18 drawn to the extent of the first through fifth appearing special technical feature and first method of use of peptide subfragments of SEQ ID Nos:2, 3, 5, 7 and 8 respectively.

Group VI-X, claim(s) 7-9, 13-15 and 16-18 drawn to the extent of the second special technical feature and first appearing use of monoclonal antibodies characterized by specifically binding the peptides of Groups I-V, respectively.

Group XI, claim 12 drawn to the second method of use of the first through fifth special technical feature.

3. The inventions listed as Groups I -XI do not relate to a single general inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, they lack the same or corresponding special technical features for the following reasons: The special technical features of Groups I-V, VI-X and XI, do not share in common the a special technical feature core structure of amino acids, antibody of identical structure and binding specificity or common us as designated in the claims. The peptides differ in amino acids, the antibodies differ in reactivity and the method of claim 12 is the second functional use of the first through fifth designated special technical feature of applicants choice.

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- 4. Applicant is advised that the reply to this requirement to be complete must include an election of the invention to be examined even though the requirement be traversed (37 CFR 1.143). For clarification, if applicants elect Group I, they elect the peptide subfragments of SEQ ID NO:2 for example. If applicants elects Group VI, they elect the monoclonal antibody reactive to a peptide subfragment of SEQ ID NO:2.
- 5. This application contains claims directed to more than one species of the generic invention. These species are deemed to lack unity of invention because they are not so linked as to form a single general inventive concept under PCT Rule 13.1.

The species are as follows: The individual species (combination/subcombination of molecules disclosed as useable together) kit components wherein the components are of the designated peptides and antibodies of Groups I-X set forth above.

Applicant is required, in reply to this action, to elect the kit species to which the claims shall be restricted if no generic claim is finally held to be allowable. The reply must also identify the claims readable on the elected species, including any claims subsequently added. An argument that a claim is allowable or that all claims are generic is considered non-responsive unless accompanied by an election.

Upon the allowance of a generic claim, applicant will be entitled to consideration of claims to additional species which are written in dependent form or otherwise include all the limitations of an allowed generic claim as provided by 37 CFR 1.141. If claims are added after the election, applicant must indicate which are readable upon the elected species. MPEP

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§ 809.02(a). For further clarification the applicant is required, regardless of the group elected, to designate the essential kit components of the claims to which the search of the claims will be initially restricted.

6. The claims are deemed to correspond to the species listed above in the following manner:

The species are those peptide subfragments and monoclonal antibodies as designated in Groups I-X as set forth above.

The following claim(s) are generic: 16-18

7. The species listed above do not relate to a single general inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, the species lack the same or corresponding special technical features for the following reasons:

The species lack the same amino acid structure and immunoreactivity as set forth above.

- 8. Applicant is reminded that upon the cancellation of claims to a non-elected invention, the inventorship must be amended in compliance with 37 CFR 1.48(b) if one or more of the currently named inventors is no longer an inventor of at least one claim remaining in the application. Any amendment of inventorship must be accompanied by a petition under 37 CFR 1.48(b) and by the fee required under 37 CFR 1.17(l).
- 9. Any inquiry of a general nature or relating to the status of this general application should be directed to the Group receptionist whose telephone number is (703) 308-0196.

Papers relating to this application may be submitted to Technology Center 1600, Group 1640 by facsimile transmission. The faxing of such papers must conform with the notice published in the Official Gazette, 1096 OG 30 (November 15, 1989). Should applicant wish to FAX a response, the current FAX number for Group 1600 is (703) 308-4242.

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Any inquiry concerning this communication or earlier communications from the examiner should be directed to Sharon L. Turner, Ph.D. whose telephone number is (703) 308-0056. The examiner can normally be reached on Monday-Friday from 8:00 AM to 4:30 PM. If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Gary Kunz, can be reached at (703) 308-4623.

Sharon L. Turner, Ph.D. August 17, 2001

CHRISTINE J. SAOUD PRIMARY EXAMINER

Trustine J. Saong